MOXIE HAND WIPES- benzalkonium chloride cloth
Skaffles Group

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Moxie Antibacterial Hand Wipes

Active Ingredient

Benzalkonium Chloride (0.13%)

Purpose

Antibacterial

DRUG FACTS

Use

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use.

Warnings

For external use only.

Do not use

- If you are allergic to any of the ingredients
- In the eyes. If contact occurs, rinse thoroughly with water.

Discontinue use if irritation and redness develop.If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Adults and children 2 yrs and over, apply to hands and allow skin to dry without wiping.
- Children under 2, ask doctor before use.

Inactive Ingredients

Water, Phenoxyethanol, Potassium Sorbate, Sodum Benzoate, Cetylpyridinium Chloride, Disodium Cocoamphodiacetate, Disodium EDTA, PEG-8 Dimethicone, Aloe Barbadensis Leaf Juice, Chamomilla Recutita (Matricacria), Flower Extract, Quaternium-52, Citric Acid, PEG075 Lanolin.

888-251-009

Principal Display Panel

Moxie

ANTIBACTERIAL HAND WIPES

Kills 99.9% of most comon gems

Moisturizes with Aloe

Unscented

6.5 in x 4.5 in (16.5 cm x 11.4 cm)

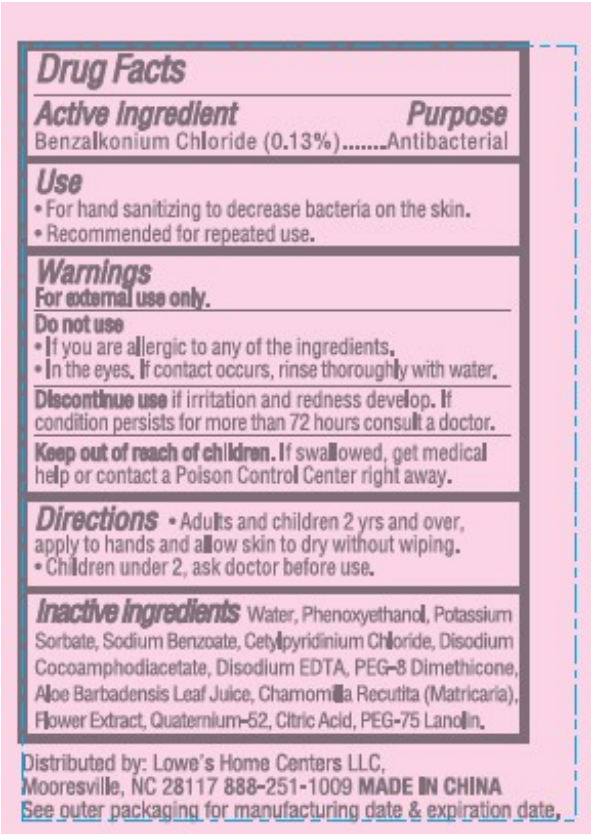
1CT

Distributed by: Lowe's Home Centers LLC,

Mooresville NC 28117 888-251-009

MADE IN CHINA

See outer packaging for manufacturing date & expiration date



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Distributed by: Lowe's Home Centers LLC,
Mooresville, NC 28117 888-251-1009 **MADE IN CHINA**
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MOXIE HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77720-035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PEG-75 LANOLIN (UNII: 09179OX7TB)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
QUATERNIUM-52 (UNII: 588EQF3H1P)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
MATRICARIA CHAMOMILLA FLOWERING TOP OIL (UNII: SA8AR2W4ER)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
WATER (UNII: 059QF0KO0R)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Product Characteristics

Color		**Score**	
Shape	RECTANGLE	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77720-035-01	1 mL in 1 POUCH; Type 0: Not a Combination Product	05/06/2024	
2	NDC:77720-035-02	50 mL in 1 POUCH; Type 0: Not a Combination Product	05/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/06/2024	

Labeler - Skaffles Group (831115642)

Revised: 5/2024 Skaffles Group